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EXAMINER

MELLER, MICHAEL V

ART UNIT

PAPER NUMBER

1655

DATE MAILED: 06/05/2006

Please find below and/or attached an Office communication concerning this application or proceeding.

DETAILED ACTION

Election/Restrictions

Applicant's election with traverse of Group I, claims 1, 2, 4-11, chondroitinase (as the enzyme), local administration and hypertrophic scars in the reply filed on 3/9/2006 is acknowledged. The traversal is on the ground(s) that the inventions are independent and that there is not an undue burden on the examiner to search both of the inventions. This is not found persuasive because the enzymes administered in the method can be used in a materially distinct method as noted by the examiner. Such enzymes can also be used to treat other conditions such as diabetes. Thus the inventions are independent and distinct from one another. Further, there is a burden on the examiner as the applicant is reminded of not only the patent search which is performed but also the extensive literature search which also must be performed on this application, thus a proper reason exists for restriction.

Thus, claims, 9, 11-13, 15-18 are withdrawn from further consideration as being drawn to non-elected subject matter.

The requirement is still deemed proper and is therefore made FINAL.

Claim Rejections - 35 USC § 102

The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless –

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

(e) the invention was described in (1) an application for patent, published under section 122(b), by another filed in the United States before the invention by the applicant for patent or (2) a patent granted on an application for patent by another filed in the United States before the invention by the applicant for patent, except that an international application filed under the treaty defined in section 351(a) shall have the effects for purposes of this subsection of an application filed in the United States only if the international application designated the United States and was published under Article 21(2) of such treaty in the English language.

(e) the invention was described in a patent granted on an application for patent by another filed in the United States before the invention thereof by the applicant for patent, or on an international application by another who has fulfilled the requirements of paragraphs (1), (2), and (4) of section 371(c) of this title before the invention thereof by the applicant for patent.

The changes made to 35 U.S.C. 102(e) by the American Inventors Protection Act of 1999 (AIPA) and the Intellectual Property and High Technology Technical Amendments Act of 2002 do not apply when the reference is a U.S. patent resulting directly or indirectly from an international application filed before November 29, 2000. Therefore, the prior art date of the reference is determined under 35 U.S.C. 102(e) prior to the amendment by the AIPA (pre-AIPA 35 U.S.C. 102(e)).

Claims 1, 2, 4-8, 10 are rejected under 35 U.S.C. 102(b) as being anticipated by WO 96/01648 (abstract, page 1, lines 20-30, page 3, lines 25-end, page 4, lines 1-10, page 29, lines 5-15, page 32, lines 20-30, page 56, lines 5-end, page 57, lines 1-10).

WO teaches that chondroitinases AC and B are well known to be from *Flavobacterium heparinium* and are known to be used to manipulate cell proliferation. WO teaches that an individual is treated after a wound (page 32, lines 20-30) and such a wound would have scarring. In fact, WO teaches that wound healing in the patients treated were even evaluated for the types of scabs that were formed which would clearly have scarring, see page 56, line 25-page 57, line 10. It is also noted that the enzymes can be administered locally, see page 29, lines 5-15.

Claims 1, 2, 6, 8, 10 are rejected under 35 U.S.C. 102(b or e) as being anticipated by Sasisekharan et al. '417 (abstract, col. 4, lines 30-40, col. 15, line 60-col. 16, line 20), WO 95/13830 (abstract, page 11, lines 25-30, the claims), or Sasisekharan et al. '430 (paragraphs 31, 63, 109, 145).

Each reference teaches that a heparinase 1, 2 or 3 from *Flavobacterium heparium* is administered locally to treat psoriasis.

Claims 1, 2, 4-8 are rejected under 35 U.S.C. 102(e) as being anticipated by Yacoby-Zeevi (col. 5, lines 23-35, col. 6, lines 1-15, 55-65, col. 12, lines 40-55, the claims).

The reference teaches that a heparinase 1, 2 or 3 from *Flavobacterium heparium* is administered to treat pulmonary fibrosis. It also teaches treating diseases that have

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angiogenesis as a pathological consequence such as cystic fibrosis, pulmonary emphysema, asthma, etc. The reference also uses chondroitinases to treat such diseases.

Claims 1, 6, 8, 10 are rejected under 35 U.S.C. 102(e) as being anticipated by Heinrichson et al. (col. 14, lines 25-40, example 1).

Heinrichson teaches topically administering heparanase to treat psoriasis.

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

Claims 1, 2, 4-8, 10 are rejected under 35 U.S.C. 103(a) as being unpatentable over Yacoby-Zeevi (col. 5, lines 23-35, col. 6, lines 1-15, 55-65, col. 12, lines 40-55, the claims) in view of Sasisekharan et al. '417 (abstract, col. 4, lines 30-40, col. 15, line 60-

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col. 16, line 20), WO 95/13830 (abstract, page 11, lines 25-30, the claims), or Sasisekharan et al. '430 (paragraphs 31, 63, 109, 145).

The reference teaches that a heparinase 1, 2 or 3 from *Flavobacterium heparium* is administered to treat pulmonary fibrosis. It also teaches treating diseases that have angiogenesis as a pathological consequence such as cystic fibrosis, pulmonary emphysema, asthma, etc. The reference also uses chondroitinases to treat such diseases.

What Yacoby does not teach is that the enzymes are administered locally.

The Sasisekharan references and the WO teach that a heparinase 1, 2 or 3 from *Flavobacterium heparium* are administered locally to treat psoriasis.

Thus since one would know to treat psoriasis locally with heparinase then one would have been motivated to treat another disease that has angiogenesis as a pathological consequence such as pulmonary emphysema locally also with a heparinase. Thus, to treat the emphysema should not be only restricted to inhalation but can also be administered locally according to the secondary references.

Claims 1, 2, 6, 8, 10 are rejected under 35 U.S.C. 103(a) as being unpatentable over Heinrikson et al. (col. 14, lines 25-40, example 1) taken with Sasisekharan et al. '417 (abstract, col. 4, lines 30-40, col. 15, line 60-col. 16, line 20), WO 95/13830

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(abstract, page 11, lines 25-30, the claims), or Sasisekharan et al. '430 (paragraphs 31, 63, 109, 145).

Heinrikson teaches topically administering heparanase to treat psoriasis.

Heinrikson does not teach that the enzymes come from the claimed microorganism.

Sasisekharan et al. '417 , Sasisekharan et al. '430 (paragraphs 31, 63, 109, 145) and WO each teach that a heparinase 1, 2 or 3 from *Flavobacterium heparium* is administered locally to treat psoriasis.

Since the secondary references each teach that heparinases from *Flavobacterium heparium* are known to be administered locally to treat psoriasis then it would have been obvious to use heparinases from *Flavobacterium heparium* instead of human since the secondary references shown that heparinases from *Flavobacterium heparium* achieve beneficial results.

Claims 1, 2, 4-8, 10 rejected under 35 U.S.C. 103(a) as being unpatentable over Sasisekharan et al. '417 (abstract, col. 4, lines 30-40, col. 15, line 60-col. 16, line 20), WO 95/13830 (abstract, page 11, lines 25-30, the claims), or Sasisekharan et al. '430 (paragraphs 31, 63, 109, 145) taken with WO 96/01648 (abstract, page 1, lines 20-30, page 3, lines 25-end, page 4, lines 1-10, page 29, lines 5-15, page 32, lines 20-30, page 56, lines 5-end, page 57, lines 1-10).

The primary references teach that a heparinase 1, 2 or 3 from *Flavobacterium heparium* is administered locally to treat psoriasis.

The primary references do not teach treating psoriasis with a chondroitinase.

WO teaches that chondroitinases AC and B are well known to be from *Flavobacterium heparinium* and are known to be used to manipulate cell proliferation. WO teaches that an individual is treated after a wound (page 32, lines 20-30) and such a wound would have scarring. In fact, WO teaches that wound healing in the patients treated were even evaluated for the types of scabs that were formed which would clearly have scarring, see page 56, line 25-page 57, line 10. It is also noted that the enzymes can be administered locally, see page 29, lines 5-15.

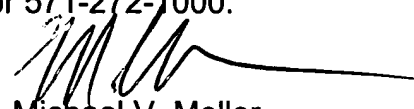
Since WO each teach that chondroitinase B from *Flavobacterium heparium* is known to be administered locally to manipulate cell proliferation then it would have been obvious to use chondroitinases from *Flavobacterium heparium* instead of the heparinase since WO shows that chondroitinases from *Flavobacterium heparium* achieve beneficial results in manipulating cell proliferation which is important in treat psoriasis since one would want to manipulate the cell proliferation to reduce the rate to control the psoriasis. One would also be motivated to use bacterial enzymes since WO clearly shows that such enzymes are well known and very beneficial which comes from microorganisms.

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Any inquiry concerning this communication or earlier communications from the examiner should be directed to Michael V. Meller whose telephone number is 571-272-0967. The examiner can normally be reached on Monday thru Thursday: 9:30am-6:00pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Terry McKelvey can be reached on 571-272-0775. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.



Michael V. Meller
Primary Examiner
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MVM